

EAST Search History

Ref #	Hits	Search Query	DBs	Default Operator	Plurals	Time Stamp
S1	1642	2,6-diisopropylphenol or propofol or propofolum	US-PGPUB; USPAT; EPO; DERWENT	OR	OFF	2006/09/19 08:13
S2	888568	injection	US-PGPUB; USPAT; EPO; DERWENT	AND	ON	2006/09/18 10:20
S3	847	S1 and S2	US-PGPUB; USPAT; EPO; DERWENT	AND	ON	2006/09/18 10:33
S4	400	polyethylene adj glycol adj "660" adj hydroxystearate or solutol	US-PGPUB; USPAT; EPO; DERWENT	OR	ON	2006/09/18 10:27
S5	794	tetrahydrofurfuryl adj alcohol adj polyethyleneglycol adj ether or glycofurol	US-PGPUB; USPAT; EPO; DERWENT	OR	ON	2006/09/18 10:28
S6	3	S3 and S4 and S5	US-PGPUB; USPAT; EPO; DERWENT	AND	ON	2006/09/18 10:28
S7	413	S1 and S2 @py<="2003"	US-PGPUB; USPAT; EPO; DERWENT	AND	ON	2006/09/18 10:34
S8	8840	bile adj salt	US-PGPUB; USPAT; EPO; DERWENT	ADJ	ON	2006/09/18 10:34
S9	56	S7 and S8	US-PGPUB; USPAT; EPO; DERWENT	ADJ	ON	2006/09/18 10:40
S10	4	S4 and S7	US-PGPUB; USPAT; EPO; DERWENT	ADJ	ON	2006/09/18 10:40
S11	1642	2,6-diisopropylphenol or propofol or propofolum	US-PGPUB; USPAT; EPO; DERWENT	OR	OFF	2006/09/18 13:15

EAST Search History

S12	888568	injection	US-PGPUB; USPAT; EPO; DERWENT	AND	ON	2006/09/18 13:15
S13	794	tetrahydrofurfuryl adj alcohol adj polyethyleneglycol adj ether or glycofurol	US-PGPUB; USPAT; EPO; DERWENT	OR	ON	2006/09/18 13:24
S14	34	S11 and S12 and S13	US-PGPUB; USPAT; EPO; DERWENT	AND	OFF	2006/09/18 13:16
S15	8	tetrahydrofurfuryl adj alcohol adj polyethyleneglycol adj ether	US-PGPUB; USPAT; EPO; DERWENT	ADJ	ON	2006/09/18 13:25
S16	1278	tetrahydrofurfuryl adj alcohol adj polyethyleneglycol adj glycofurol or tetraglycol or methoxy adj PEG	US-PGPUB; USPAT; EPO; DERWENT	OR	ON	2006/09/18 13:31
S17	18	S11 and S12 and S16	US-PGPUB; USPAT; EPO; DERWENT	OR	ON	2006/09/18 13:27
S18	1102762	tetrahydrofurfuryl adj alcohol adj polyethyleneglycol adj glycofurol or tetraglycol or methoxy adj PEG or tetrahydrofurfuryl alcohol PEG ether	US-PGPUB; USPAT; EPO; DERWENT	OR	ON	2006/09/18 13:31
S19	1280	tetrahydrofurfuryl adj alcohol adj polyethyleneglycol adj glycofurol or tetraglycol or methoxy adj PEG or tetrahydrofurfuryl adj alcohol adj PEG adj ether	US-PGPUB; USPAT; EPO; DERWENT	OR	ON	2006/09/18 13:31
S20	113	diprivan	US-PGPUB; USPAT; EPO; DERWENT	AND	ON	2006/09/19 08:17
S21	1665	diprivan or propofol or 2, 6-diisopropylphenol or propofolum	US-PGPUB; USPAT; EPO; DERWENT	OR	ON	2006/09/19 08:17
S22	1280	tetrahydrofurfuryl adj alcohol adj polyethyleneglycol adj glycofurol or tetraglycol or methoxy adj PEG or tetrahydrofurfuryl adj alcohol adj PEG adj ether	US-PGPUB; USPAT; EPO; DERWENT	OR	ON	2006/09/19 08:17

EAST Search History

S23	30	S21 and S22	US-PGPUB; USPAT; EPO; DERWENT	OR	ON	2006/09/19 08:18
S24	30	S21 and S22	US-PGPUB; USPAT; EPO; DERWENT	AND	ON	2006/09/19 08:18

FILE 'HOME' ENTERED AT 10:03:00 ON 19 SEP 2006

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=> s 2,6-diisopropylphenol or propofol or propofolum
L1 39816 2,6-DIISOPROPYLPHENOL OR PROPOFOL OR PROPOFOLUM

=> s injection or microemulsion
L2 1617420 INJECTION OR MICROEMULSION

=> s l1 and l2
L3 4299 L1 AND L2

=> s polyethylene(w)glycol(w)660(w)hydroxystearate or solutol
L4 362 POLYETHYLENE(W) GLYCOL(W) 660(W) HYDROXYSTEARATE OR SOLUTOL

=> s tetrahydrofurfuryl(w)alcohol(w)polyethyleneglycol(w)ether or glycofurol or
tetraglycol or methoxy(w)PEG
L5 786 TETRAHYDROFURFURYL(W) ALCOHOL(W) POLYETHYLENEGLYCOL(W) ETHER OR
GLYCOFUROL OR TETRAGLYCOL OR METHOXY(W) PEG

=> s l3 and l4 and l5
L6 2 L3 AND L4 AND L5

=> d ti au abs so py 1-2

L6 ANSWER 1 OF 2 CAPLUS COPYRIGHT 2006 ACS on STN
TI Formulations containing propofol for anesthetic use
IN Bell, Alan R.; Cochrane, Fenella; O'Connor, Geoffrey N.; Rowe, James S.
AB A formulation for anesthetic use is described. The formulation contains
propofol, and may be used to induce and/or maintain anesthesia or
sedation in a vertebrae. The formulation addnl. contains a solvent or a
combination of solvents and is suitable for mixing with commonly used
infusion fluids prior to injection to a patient. The
formulation may be terminally sterilized using moist heat in order to
assure sterility, and contains no lipid, thereby avoiding complications
associated with administration over prolonged periods of time, or to patients
with disorders of fat metabolism For example, a solution was formulated
containing
propofol 1, glycofurol 20, Solutol HS15 10,
benzyl alc. 2, ethanol 2 % weight/volume, and water for injection to
100 %.
SO U.S. Pat. Appl. Publ., 12 pp.
CODEN: USXXCO
PY 2005
2005
2004
2004

L6 ANSWER 2 OF 2 CAPLUS COPYRIGHT 2006 ACS on STN
TI Injectable 2,6-diisopropylphenol-containing
anesthetic composition and methods
IN Jee, Ung-kil
AB An injectable anesthetic composition in a microemulsion phase is
disclosed. The composition contains 2,6-
diisopropylphenol as the active ingredient, together with
polyethylene glycol 660
hydroxystearate, tetrahydrofurfuryl alc.
polyethyleneglycol ether, and an aqueous medium. Methods of
making and using the injectable anesthetic composition are also disclosed.
SO U.S. Pat. Appl. Publ., 9 pp.
CODEN: USXXCO
PY 2004
2004
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2004
2005
2005

=> s l3 and l5

L7 2 L3 AND L5

=> d ti au so py 1-2

L7 ANSWER 1 OF 2 CAPLUS COPYRIGHT 2006 ACS on STN
TI Formulations containing propofol for anesthetic use
IN Bell, Alan R.; Cochrane, Fenella; O'Connor, Geoffrey N.; Rowe, James S.
SO U.S. Pat. Appl. Publ., 12 pp.
CODEN: USXXCO
PY 2005
2005
2004
2004

L7 ANSWER 2 OF 2 CAPLUS COPYRIGHT 2006 ACS on STN
TI Injectable 2,6-diisopropylphenol-containing
anesthetic composition and methods
IN Jee, Ung-kil
SO U.S. Pat. Appl. Publ., 9 pp.
CODEN: USXXCO
PY 2004
2004
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2004
2005
2005

=> s cellulose

L8 466581 CELLULOSE

=> s l3 and l8

L9 3 L3 AND L8

=> s ti au abs py so 1-3

L10 0 TI AU ABS PY SO 1-3

=> d ti au abs py so 1-3

L10 HAS NO ANSWERS

'1-3 ' IS NOT A VALID SEARCH STATUS KEYWORD
Search status keywords:

NONE ---- Display only the number of postings.
STATUS -- Display statistics of the search.
ENTER SEARCH STATUS OPTION (NONE), STATUS, OR ?:19
'L45' IS NOT A VALID SEARCH STATUS KEYWORD
Search status keywords:
NONE ---- Display only the number of postings.
STATUS -- Display statistics of the search.
ENTER SEARCH STATUS OPTION (NONE), STATUS, OR ?:none
L10 0 SEA TI AU ABS PY SO 1-3

=> d 19 ti au abs py so 1-3

L9 ANSWER 1 OF 3 CAPLUS COPYRIGHT 2006 ACS on STN
TI Novel encochleation methods, cochleates and methods of use
IN Mannino, Raphael J.; Gould-Fogerite, Susan; Krause-Elsmore, Sara L.;
 Delmarre, David; Lu, Ruying
AB The invention generally relates to cochleate drug delivery vehicles.
 Disclose are novel methods for making cochleates and cochleate compns.
 that include introducing a cargo moiety to a liposome in the presence of a
 solvent. Also disclosed are cochleates and cochleate compns. that include
 an aggregation inhibitor, and optionally, a cargo moiety. Addnl., anhydrous
 cochleates that include a protonized cargo moiety, a divalent metal cation
 and a neg. charge lipid are disclosed. Methods of using the cochleate
 compns. of the invention, including methods of administration, are also
 disclosed.
PY 2004
 2005
 2005
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 2006
SO PCT Int. Appl., 195 pp.
 CODEN: PIXXD2

L9 ANSWER 2 OF 3 CAPLUS COPYRIGHT 2006 ACS on STN
TI Injectable 2,6-diisopropylphenol-containing
 anesthetic composition and methods
IN Jee, Ung-kil
AB An injectable anesthetic composition in a microemulsion phase is
 disclosed. The composition contains 2,6-
 diisopropylphenol as the active ingredient, together with
 polyethylene glycol 660 hydroxystearate, tetrahydrofurfuryl alc.
 polyethyleneglycol ether, and an aqueous medium. Methods of making and using
 the injectable anesthetic composition are also disclosed.
PY 2004
 2004
 2004
 2004
 2005
 2005
SO U.S. Pat. Appl. Publ., 9 pp.
 CODEN: USXXCO

L9 ANSWER 3 OF 3 CAPLUS COPYRIGHT 2006 ACS on STN
TI Biodegradable injectable implants and related methods of manufacture and
 use
IN Caseres, Crisofo Peralta; D'Lagarde, Daniel Leon
AB This invention is directed to the field of medical implants, and more
 specifically to biodegradable injectable implants and their methods of
 manufacture and use. The injectable implants disclosed herein comprise
 glycolic acid and bio-compatible/bio-absorbable polymeric particles containing
 a polymer of lactic acid. The particles are small enough to be injected
 through a needle but large enough to avoid engulfment by macrophages. The
 injectables of this invention may be in a pre-activated solid form or an

activated form (e.g., injectable suspension or emulsion). For example, a lyophilized composition was prepared containing glycolic acid 0.07 mg, poly(lactic acid) spheres 200.0 mg, hydroxypropyl Me cellulose 118.33 mg, D-mannitol 170.0 mg, pH stabilizer (phosphate buffer) 0.50 mg, and surfactant (Tween 80) 1.20 mg. The composition was activated extemporaneously with 5.5 mL water to obtain an injectable preparation

PY 2003
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SO PCT Int. Appl., 60 pp.
CODEN: PIXXD2

=> s pH(w)regulator

L11 1080 PH(W) REGULATOR

=> s l3 and l11

L12 0 L3 AND L11

=> s acetate

L13 884590 ACETATE

=> s l3 and l13

L14 31 L3 AND L13

=> d ti au abs so py 1-5

L14 ANSWER 1 OF 31 CAPLUS COPYRIGHT 2006 ACS on STN

TI Lipid-based dispersions for drug delivery

IN Hu, Ning; Jensen, Gerard M.; Yang, Stephanie; Su-ming, Chiang

AB The invention provides lipid-based dispersion comprising comprising, phosphatidylcholine, an anionic phospholipid, up to 1% cholesterol by weight of total lipids, and a therapeutic agent, wherein the mean particle size measured by dynamic light scattering is <100 nm. The invention also provides pharmaceutical comps. comprising such a dispersion as well as methods of producing a therapeutic effect in a mammal comprising administering an effective amount of such a dispersion. Soy-phosphatidylcholine, DSPG, and propofol were dissolved in a 1:1 mixture of methanol and chloroform at a molar ratio of Soy-PC:DSPG of 1:0.4 and a weight ratio of (Soy-PC + DSPG):propofol of 10:1. Solvents were removed by evaporation and the films were then hydrated in 9% sucrose at desired drug concns. and sonicated to form liposomes.

SO PCT Int. Appl., 31 pp.
CODEN: PIXXD2

PY 2005
2006
2005
2005
2005

L14 ANSWER 2 OF 31 CAPLUS COPYRIGHT 2006 ACS on STN

TI Propofol formulation containing reduced oil and surfactants

IN Desai, Neil P.; Yang, Andrew; De, Tapas; Ci, Sherry Xiaopei; Soon-Shiong, Patrick

AB Sterile, stable pharmaceutical formulations of emulsions of neat propofol or propofol dissolved in a solvent and containing no preservative are provided that comprise optimal amts. of surfactants such as lecithin and solvent such as soybean oil, with a suitable pH range

to prevent significant growth of microorganisms for at least 24 h after extrinsic contamination. The lower amount of oil or absence (oil) in the formulation also allows chronic sedation over extended periods of time with a reduced chance of lipid overload in the blood. Formulations with the following general ranges of components for such propofol compns. were prepared as follows: propofol 0.5-5; human serum albumin 0.01-3; glycerol 2.25; water for injection qs to 100%.

SO U.S. Pat. Appl. Publ., 13 pp.

CODEN: USXXCO

PY 2004

2004

2005

L14 ANSWER 3 OF 31 CAPLUS COPYRIGHT 2006 ACS on STN

TI Propofol with cysteine

IN Tang, Hua; Chen, Hongming; Almarsson, Orn

AB The present invention relates to pharmaceutical compns. comprising 2,6-diisopropylphenol (propofol).

Compns. of the present invention comprise aqueous and non-aqueous compns. of propofol and cysteine or a salt thereof. The propofol containing compns. are preferably sterile and are parenterally administered to any animal, including humans.

SO PCT Int. Appl., 23 pp.

CODEN: PIXXD2

PY 2004

2004

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L14 ANSWER 4 OF 31 CAPLUS COPYRIGHT 2006 ACS on STN

TI Substituted phenol compounds useful for anesthesia and sedation

IN Jenkins, Thomas E.; Ji, Yu-Hua; Wu, Huiwei; Bolton, Jennifer

AB Substituted phenol compds. and pharmaceutical compns. containing them, e.g., injections, which are useful for inducing or maintaining anesthesia or sedation in a mammal are provided. For example, 4-methoxycarbonyl-2,6-diisopropylphenol, prepared by the reaction of carbon tetrachloride, 2,6-diisopropylphenol and methanol, was used for the synthesis of 4-(2-methoxycarbonylethyl)-2,6-diisopropylphenol by its reaction with Et acetate.

SO PCT Int. Appl., 41 pp.

CODEN: PIXXD2

PY 2003

2003

2003

2003

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2004

2005

L14 ANSWER 5 OF 31 CAPLUS COPYRIGHT 2006 ACS on STN

TI Novel pharmaceuticals comprising drug conjugates with polypeptide carriers

IN Picariello, Thomas

AB A pharmaceutical composition comprising a polypeptide and an active agent attached to said polypeptide is disclosed.

SO PCT Int. Appl., 2059 pp.

CODEN: PIXXD2

PY 2003

2003

2003

2003
2006
2004
2006

=> s diprivan or propofol or 2,6-diisopropylphenol or propofolum
L15 39943 DIPRIVAN OR PROPOFOL OR 2,6-DIISOPROPYLPHENOL OR PROPOFOLUM

=> d his

(FILE 'HOME' ENTERED AT 10:03:00 ON 19 SEP 2006)

FILE 'CAPLUS, MEDLINE, BIOSIS, EMBASE' ENTERED AT 10:03:28 ON 19 SEP 2006
L1 39816 S 2,6-DIISOPROPYLPHENOL OR PROPOFOL OR PROPOFOLUM
L2 1617420 S INJECTION OR MICROEMULSION
L3 4299 S L1 AND L2
L4 362 S POLYETHYLENE(W) GLYCOL(W) 660(W) HYDROXYSTEARATE OR SOLUTOL
L5 786 S TETRAHYDROFURFURYL(W) ALCOHOL(W) POLYETHYLENEGLYCOL(W) ETHER OR
L6 2 S L3 AND L4 AND L5
L7 2 S L3 AND L5
L8 466581 S CELLULOSE
L9 3 S L3 AND L8
L10 0 S TI AU ABS PY SO 1-3
L11 1080 S PH(W) REGULATOR
L12 0 S L3 AND L11
L13 884590 S ACETATE
L14 31 S L3 AND L13
L15 39943 S DIPRIVAN OR PROPOFOL OR 2,6-DIISOPROPYLPHENOL OR PROPOFOLUM

=> s l15 and l2

L16 4308 L15 AND L2

=> s l16 and l4 and l5

L17 2 L16 AND L4 AND L5

=> s l16 and l11

L18 0 L16 AND L11

=> s l16 and l8

L19 3 L16 AND L8

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